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		Serial Number-Kind Code <sup>2</sup> (if known)			

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Substitute for form 1449/PTO			<b>Complete If Known</b>	
<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>			<b>Application Number</b>	10/567,472
			<b>Filing Date</b>	Feb. 7, 2006
Date Submitted: September 19, 2007 (use as many sheets as necessary)			<b>First Named Inventor</b>	DOBROVOLNY, Petr
			<b>Art Unit</b>	1625
Sheet 2 of 3			<b>Examiner Name</b>	Aulakh, Charanjit
			<b>Attorney Docket Number</b>	087329-0108

FOREIGN PATENT DOCUMENTS						
Examiner Initials*	Cite No. <sup>1</sup>	Foreign Patent Document Country Code <sup>3</sup> Number <sup>4</sup> Kind Code <sup>5</sup> (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Documents	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>
CA	A23	EP74256	11-20-1986	Miyasaka et al.		
CA	A24	EP74770	03-23-1983	Miyasaka et al.		
CA	A25	EP88642	09-14-1983	Miyasaka et al.		
CA	A26	EP154583	09-11-1988	Funck et al.		
CA	A27	EP154584	02-03-1988	Pressaco et al.		
CA	A28	EP51289	04-09-1986	Gelert		
CA	A29	WO96/31513	10-10-1996	Henegar et al.		
CA	A30	WO2004/100897	11-25-2004	Lin et al.		
CA	A31	WO2005/019223	03-03-2005	Dobrovoly		
CA	A32	WO2005/058910	06-30-2005	Dobrovoly		

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.) date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>6</sup>
CA	A33	"Combination of Irinotecan (CPT-11) and 5-Fluorouracil with an analysis of cellular determinants of drug activity," PAVILLARD et al. Biochemical Pharmacology, vol. 56: 1315-1322, 1998.*	
CA	A34	"Clinical advances with topoisomerase I inhibitors in gastrointestinal malignancies," ARMAND, Jean-Pierre et al. Anti-Cancer Drugs 10 (Suppl. 1): S5-S12 (1999).	
CA	A35	"Phase I/II study of escalating dose of CPT-11 in combination with LV5FU2 ("De Gramont" regimen) every 2 weeks in the treatment of colorectal cancer (CRC) after 5-FU failure," DUCREUX, M. et al., Abstract 823, Proc. of Amer. Soc. Clin. Oncol. 16:234a (1997).	
CA	A36	"Phase I study of a weekly schedule of irinotecan (CPT-11), high-dose folinic acid (FA) and 5-fluorouracil (5-FU) as first line chemotherapy (CT) in metastatic colorectal cancer: Final results," VANHOEFER, U. et al., Abstract 779, Proc. of Amer. Soc. Clin. Oncol. 17:202a (1998).	
CA	A37	"Irinotecan (CPT-11) in the treatment of gastrointestinal cancers," NISHIYAMA, M. Japanese J. Chemotherapy 46(8):292-296 (1998).	
CA	A38	"CPT-11 (Irinotecan) and 5-Fluorouracil: a Promising Combination for Therapy of Colorectal Cancer," SALTZ, L., et al. European J. Cancer 32A(Suppl. 3):S24-S31 (1996).	

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CA	A39	"Phase I/II study of CPT-11 in combination with LV5FU2 (De Gramont-Regimen) every 2 weeks for the treatment of colorectal cancer (CRC) after 5-FU failure," SEITZ, J.F. et al. Abstract 261, Annals of Oncology 9 (Suppl. 2):68 (1998).		
CA	A40	"Phase I study of a weekly schedule of irinotecan (CPT-11) in combination with high-dose folinic acid and 5-fluorouracil as first line chemotherapy in patients with advanced colorectal cancer," VANHOEFER, U. et al. Abstract 967, Proc. of Amer. Soc. Clin. Oncol. 16:272a (1997).		
CA	A41	"Synthesis and Antitumor Activity of 20(S)-Camptothecin Derivatives. A-Ring-Substituted 7-Ethylcamptothecins and their E-Ring-Modified Water-Soluble Derivatives," YAEGASHI et al. Chemical & Pharmaceutical Bulletin. Vol. 42. No. 12: 2518-2525(1994).		
CA	A42	"Chemical Modification of an Antitumor Camptothecin: Synthesis and Antitumor Activity of 7-C-Substituted Camptothecins," SAWADA et al. Chemical & Pharmaceutical Bulletin. Vol. 39. No. 10: 2574-2580 (1991).		
CA	A43	"Synthesis and Antitumor Activity of A-Ring or E-Lactone Modified Water-Soluble Prodrugs of 20(S)-Camptothecin, Including Development of Irinotecan Hydrochloride Trihydrate," SAWADA et al. Current Pharmaceutical Design. Vol. 1 No. 1: 113-132 (1995).		
CA	A44	Photodegradation reactions of CPT-II, a derivative of camptothecin. I: chemical structure of main degradation products in an aqueous solution," AKIMOTO et al. Drug Stability. Vol. 1 No. 2.: 118-122 (1996).		
CA	A45	"An Efficient Conversion of Camptothecin to 10-Hydroxycamptothecin," WOOD et al. The Journal of Organic Chemistry. Vol. 60. No. 17: 5739-5740 (1995).		
CA	A46	"Synthesis and Antitumor Activity of 20(S)-Camptothecin Derivatives: Carbamate-Linked, Water-Soluble Derivatives of 7-Ethyl-10-hydroxycamptothecin," SAWADA, et al. Chemical & Pharmaceutical Bulletin, Vol. 39. No. 6: 1446-1454 (1991).		
CA	A47	"Synthesis and Antitumor Activity of 20(S)-Camptothecin Derivatives: A-Ring Modified and 7, 10-Disubstituted Camptothecins," SAWADA et al. Chemical & Pharmaceutical Bulletin, Vol. 39. No. 12: 3183-3188 (1991).		

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